PTO/SB/08a (08-03)
Approved for use through 97/31/2006, ONB 0651-0201
U.S. Patient and Trademark Office, U.S. DEPARTMENT OF COMMERCE to a collection of information unfers it contains a valid OMS control number. Under the Paperwork Reduction Act of 1995, no persons are req

INFORMATION DISCLOSURE
STATEMENT BY APPLICANT
(Not for submission under 37 CFR 1.99)

Application Number		10563846			
Filing Date		2006-01-05			
First Named Inventor Thom		as FALCK, ET AL			
Art Unit		2617			
Examiner Name					
Attorney Docket Number		DE030235			

					U.S.	PATENTS			Remove		
Examiner Initial* Cite No Patent Number			Kind Code ¹ Issue Date Name of Patentee or Applican of cited Document			Pages,Columns,Lines where Relevant Passages or Relevant Figures Appear					
	1										
If you wisl	h to a	dd additional U.S. Pate	nt citatio	n inform	ation pl	lease click the	Add button.		Add		
			U.S.P	ATENT	APPLI	CATION PUB	LICATIONS		Remove		
Examiner Cite Publication Number		Kind Code ¹	Publica Date	ition	of cited Decument Releva			s,Columns,Lines where vant Passages or Relevant es Appear			
	1	20020196378	A1	2002-12	:-26	SLOBODIN E	ΓAL				
	2	20030017846	A1	2003-01	1-23	ESTEVEZ ET	AL				
If you wis	h to a	dd additional U.S. Publi	shed Ap	plication	citatio	n information p	olease click the Ad	d button	Add		
				FOREIG	SN PAT	ENT DOCUM	ENTS		Remove		
Examiner Initial*			Kind Code ⁴	Publication Date	Name of Patente Applicant of cited Document	e or V	vhere Rel	or Relevant	Τ.		
	1	0233687	wo		A2	2002-04-25	DVIR ET AL				С
	2	1244303	₽		A2	2002-09-25	VICARI				С

INFORMATION DISCLOSURE STATEMENT BY APPLICANT (Not for submission under 37 CFR 1.99)

oplication Number		10563846				
ling Date		2006-01-05				
st Named Inventor Thom		as FALCK, ET AL				
t Unit		2617				
caminer Name						
towari Danket Montes		DE02022E				

If you wisl	h to a	dd add	ditional Foreign	n Patent	Docum	ent citat	tion inf	formation	please	click the Ad	d buttor	A	dd		
				N	ION-PA	TENT L	ITERA	ATURE D	OCUME	NTS		Ren	nove		
Examinor Cite Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, serial, symposium, catalog, etc), date, pages(s), volume-issue number(s), publisher, ctly anotic country where publisher.								n T5							
	1														
If you wis	h to a	dd add	fitional non-p	atent liter	rature do	ocument	t citatio	on informa	ation ple	ase click th	ne Add b	utton	Add		
						EXAMI	INER S	SIGNATU	RE						
Examiner	Signa	ture							- 1	ate Consid	dered				
			reference co											rough	a

1 See Kind Codes of USPTO Patent Documents at www.ISPTO.QQU/or MPEP 90104. 2 Enter office that issued the document, by the hor-letter code (WIPO Standard ST3.) 2 for Junganese patent coursets, the included not not be year of the insight of the Emperor may precede the serial runber of the patent of being of the Emperor may precede the serial runber of the patent of being of the Emperor may precede the serial runber of the patent of t

INFORMATION DISCLOSURE STATEMENT BY APPLICANT (Not for submission under 37 CFR 1.99)

Application Number		10563846			
Filing Date		2006-01-05			
First Named Inventor	Thom	as FALCK, ET AL			
Art Unit		2617			
Examiner Name					
Attorney Docket Number		DE030235			

CERTIFICATION STATEMENT

Please see 37	CFR 1.97	and 1.98 to	make the ap	propriate selection(s):
---------------	----------	-------------	-------------	-------------------------

That each item of information contained in the information disclosure statement was first cited in any communication from a foreign patent office in a counterpart foreign application not more than three months prior to the filling of the information disclosure statement. See 37 CFR 1.97(e/11).

OR

That no item of information contained in the information disclosure statement was cited in a communication from a foreign patent office in a counterpart foreign application, and, to the knowledge of the person signing the certification after making reasonable inquity, no item of information contained in the information disclosure statement was known to any individual designated in 37 CFR 156(c) more than three months prior to the filing of the information disclosure statement. See 37 CFR 157(4)(c).

- See attached certification statement.
- | Fee set forth in 37 CFR 1.17 (p) has been submitted herewith.
- □ None

SIGNATURE

A signature of the applicant or representative is required in accordance with CFR 1.33, 10.18. Please see CFR 1.4(d) for the form of the signature.

Torri or the signature.			
Signature	/Aaron M. Waxler/	Date (YYYY-MM-DD)	2007-02-08
Name/Print	Aaron Waxler	Registration Number	48.027

This collection of information is required by 3T CFR 1.97 and 1.98. The information is required to obtain or retain a benefit by the public which is for life railed by the USPTO to process) and application. Confidentiality is governed by \$5 U.S. C. 12.04 and 3T CFR 1.14. This collection is estimated to take 1 hour to complete, including gathering, preparing and submitting the completed application from the USPTO. Time will vary depending upon the individual case. Any comments on the amount of time you require to complete his form and/or suggestions for reducing this burden, should be sent to the Chief Information Officer, U.S. Patent and Tradenar's Office, U.S. operatment of Commence, P. 0. Box 1450, Alexandrin, V.S. 2231-1450. DIN OTS CRID FEES OR COMPLETED FORMS TO THIS ADDRESS. SEND TO: Commissioner for Patents, P.O. Box 1450, Alexandria, V.S. 2231-1450.

Privacy Act Statement

The Privacy Act of 1974 (P. L. 93-579) requires that you be given certain information in connection with your submission of the stackhold from related to a patient application or patient. Accordingly, pursuant to the requirements of the Act, places be advised that (1) the general authority for the collection of this information is 35 U.S.C. 2(b)(2); (2) familishing of the information solicided is couldrain; and (3) the primoral pursuance for which the information is used by the U.S. Patient and Trademan Coffice is to process and/or examine your submission related to a patient agricultant or patient. If you do not furnish the requested process and/or examine your submission related to a patient agricultant or patient. If you do not furnish the requested results of the patient of the patient and the patient of the patient

The information provided by you in this form will be subject to the following routine uses:

- The information on this form will be treated confidentially to the extent allowed under the Freedom of Information Act (5 U.S.C. 552) and the Privacy Act (5 U.S.C. 552a). Records from this system of records may be disclosed to the Department of Justice to determine whether the Freedom of Information Act requires disclosure of these record s.
 - A record from this system of records may be disclosed, as a routine use, in the course of presenting evidence to a court, magistrate, or administrative tribunal, including disclosures to opposing counsel in the course of settlement negotiation.
 - A record in this system of records may be disclosed, as a routine use, to a Member of Congress submitting a request involving an individual, to whom the record pertains, when the individual has requested assistance from the Member with respect to the subject matter of the record.
 - A record in this system of records may be disclosed, as a routine use, to a contractor of the Agency having need for the information in order to perform a contract. Recipients of information shall be required to comply with the requirements of the Privacy Act of 1974, as amended, pursuant to 5 U.S.C. 552(m).
 - A record related to an International Application filed under the Patent Cooperation Treaty in this system of records
 may be disclosed, as a routine use, to the International Bureau of the World Intellectual Property Organization, pursuant
 to the Patent Cooperation Treaty.
 - A record in this system of records may be disclosed, as a routine use, to another federal agency for purposes of National Security review (35 U.S.C. 181) and for review pursuant to the Atomic Energy Act (42 U.S.C. 218(c)).
 - 7. A record from this system of records may be disclosed, as a routine use, to the Administrator, General Services, or hisher designed, uturing an insection of records conducted by GSA as part of that agency's responsibility to recommend improvements in records management practices and programs, under authority of 4d U.S.C. 2904 and 2905. Such disclosure shall be made in accordance with the GSA requisions governing inseption of records for this purpose, and any other relevant (i.e., GSA or Commerce) directive. Such disclosure shall not be used to make determinations about individuals.
- A record from this system of records may be disclosed, as a routine use, to the public after either publication of the
 application pursuant to 35 U.S.C. 12(2) or issuance of a patent pursuant to 35 U.S.C. 151. Further, a record may be
 disclosed, subject to the limitations of 37 CFR 1.14, as a routine use, to the public if the record was filled in application
 which became abandoned or in which the proceedings were terminated and which application is referenced by either a
 published application, an application open to public inspections or as issued patent.
 - A record from this system of records may be disclosed, as a routine use, to a Federal, State, or local law enforcement agency, if the USPTO becomes aware of a violation or potential violation of law or regulation.